The SPEAKER pro tempore. Pursuant to House Resolution 214, the previous question is ordered on the amendment offered by the gentleman from Virginia (Mr. SCOTT).

The question is on the amendment offered by the gentleman from Virginia (Mr. SCOTT).

The amendment was agreed to.

AMENDMENT IN THE NATURE OF A SUBSTITUTE OFFERED BY MR. GREENWOOD

Mr. GREENWOOD. Mr. Speaker, I offer an amendment in the nature of a substitute.

The SPEAKER pro tempore. The Clerk will designate the amendment in the nature of a substitute.

The text of the amendment in the nature of a substitute is as follows:

Amendment in the nature of a substitute printed in House Report 107–172 offered by Mr. GREENWOOD:

Strike all after the enacting clause and insert the following:

SECTION 1. SHORT TITLE.

This Act may be cited as the ''Cloning Prohibition Act of 2001''.

SEC. 2. PROHIBITION AGAINST HUMAN CLONING.

(a) IN GENERAL.—The Federal Food, Drug, and Cosmetic Act (21 U.S.C. 301 et seq.) is amended by adding at the end the following:

''CHAPTER X—HUMAN CLONING

''PROHIBITION AGAINST HUMAN CLONING

''SEC. 1001. (a) NUCLEAR TRANSFER TECHNOLOGY.—

''(1) IN GENERAL.—It shall be unlawful for any person—

''(A) to use or attempt to use human somatic cell nuclear transfer technology, or the product of such technology, to initiate a pregnancy or with the intent to initiate a pregnancy; or

''(B) to ship, mail, transport, or receive the product of such technology knowing that the product is intended to be used to initiate a pregnancy.

''(2) DEFINITION.—For purposes of this section, the term 'human somatic cell nuclear transfer technology' means transferring the nuclear material of a human somatic cell into an egg cell from which the nuclear material has been removed or rendered inert.

''(b) RULE OF CONSTRUCTION.—This section may not be construed as applying to any of the following:

''(1) The use of somatic cell nuclear transfer technology to clone molecules, DNA, cells, or tissues.

''(2) The use of mitochondrial, cytoplasmic, or gene therapy.

''(3) The use of in vitro fertilization, the administration of fertility-enhancing drugs, or the use of other medical procedures (excluding those using human somatic cell nuclear transfer or the product thereof) to assist a woman in becoming or remaining pregnant

''(4) The use of somatic cell nuclear transfer technology to clone or otherwise create animals other than humans.

''(5) Any other activity (including biomedical, microbiological, or agricultural research or practices) not expressly prohibited in subsection (a).

''(c) REGISTRATION.—

''(1) IN GENERAL.—Each individual who intends to perform human somatic cell nuclear transfer technology shall, prior to first performing such technology, register with the Secretary his or her name and place of business (except that, in the case of an individual who performed such technology before the date of the enactment of the Cloning Prohibition Act of 2001, the individual shall so register not later than 60 days after such date). The Secretary may by regulation require that the registration provide additional information regarding the identity and business locations of the individual, and information on the training and experience of the individual regarding the performance of such technology.

''(2) ATTESTATION.—A registration under paragraph (1) shall include a statement, signed by the individual submitting the registration, declaring that the individual is aware of the prohibitions described in subsection (a) and will not engage in any violation of such subsection.

''(3) CONFIDENTIALITY.—Information provided in a registration under paragraph (1) shall not be disclosed to the public by the Secretary except to the extent that—

''(A) the individual submitting the registration has in writing authorized the disclosure; or

''(B) the disclosure does not identify such individual or any place of business of the individual.

''(d) PREEMPTION OF STATE LAW.—This section supersedes any State or local law that—

''(1) establishes prohibitions, requirements, or authorizations regarding human somatic cell nuclear transfer technology that are different than, or in addition to, those established in subsection (a) or (c); or

''(2) with respect to humans, prohibits or restricts research regarding or practices constituting—

''(A) somatic cell nuclear transfer;

''(B) mitochondrial or cytoplasmic therapy; or

''(C) the cloning of molecules, DNA, cells, tissues, or organs;

except that this subsection does not apply to any State or local law that was in effect as of the day before the date of the enactment of the Cloning Prohibition Act of 2001.

''(e) RIGHT OF ACTION.—This section may not be construed as establishing any private right of action.

''(f) DEFINITION.—For purposes of this section, the term 'person' includes governmental entities.

''(g) SUNSET.—This section and section 301(bb) do not apply to any activity described in subsection (a) that occurs on or after the expiration of the 10-year period beginning on the date of the enactment of the Cloning Prohibition Act of 2001.''.

(b) PROHIBITED ACTS.—

(1) IN GENERAL.—Section 301 of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 331) is amended by adding at the end the following:

''(bb) The violation of section 1001(a), or the failure to register in accordance with section 1001(c).''.

(2) CRIMINAL PENALTY.—Section 303(b) of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 333(b)) is amended by adding at the end the following:

''(7) Notwithstanding subsection (a), any person who violates section 301(bb) shall be imprisoned not more than 10 years or fined in accordance with title 18, United States Code, or both.''.

(3) CIVIL PENALTY.—Section 303 of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 333) is amended by adding at the end the following:

''(h)(1) Any person who violates section 301(bb) shall be liable to the United States for a civil penalty in an amount not to exceed the greater of—

''(A) $1,000,000; or

''(B) an amount equal to the amount of any gross pecuniary gain derived from such violation multiplied by 2.

''(2) Paragraphs (3) through (5) of subsection (g) apply with respect to a civil penalty under paragraph (1) of this subsection to the same extent and in the same manner as such paragraphs (3) through (5) apply with respect to a civil penalty under paragraph (1) or (2) of subsection (g).''.

(4) FORFEITURE.—Section 303 of the Federal Food, Drug, and Cosmetic Act, as amended by paragraph (3), is amended by adding at the end the following:

''(i) Any property, real or personal, derived from or used to commit a violation of section 301(bb), or any property traceable to such property, shall be subject to forfeiture to the United States.''.

SEC. 3. STUDY BY INSTITUTE OF MEDICINE.

(a) IN GENERAL.—The Secretary of Health and Human Services (referred to in this section as the ''Secretary'') shall request the Institute of Medicine to enter into an agreement with the Secretary under which such Institute conducts a study to—

(1) review the current state of knowledge about the biological properties of stem cells obtained from embryos, fetal tissues, and adult tissues;

(2) evaluate the current state of knowledge about biological differences among stem cells obtained from embryos, fetal tissues, and adult tissues and the consequences for research and medicine; and

(3) assess what is currently known about the ability of stem cells to generate neurons, heart, kidney, blood, liver and other tissues and the potential clinical uses of these tissues.

(b) OTHER ENTITIES.—If the Institute of Medicine declines to conduct the study described in subsection (a), the Secretary shall enter into an agreement with another appropriate public or nonprofit private entity to conduct the study.

(c) REPORT.—The Secretary shall ensure that, not later than three years after the date of the enactment of this Act, the study required in subsection (a) is completed and a report describing the findings made in the study is submitted to the Committee on Energy and Commerce in the House of Representatives and the Committee on Health, Education, Labor, and Pensions in the Senate.

The SPEAKER pro tempore. Pursuant to House Resolution 214, the gentleman from Pennsylvania (Mr. GREENWOOD) and the gentleman from Wisconsin (Mr. SENSENBRENNER) each will control 30 minutes.

PARLIAMENTARY INQUIRY

Mr. GREENWOOD. Mr. Speaker, I have a parliamentary inquiry.

The SPEAKER pro tempore. The gentleman will state it.

Mr. GREENWOOD. Would it be appropriate for me or permissible under the rules for me to yield 15 minutes of my time to the gentleman from Florida (Mr. DEUTSCH)?

The SPEAKER pro tempore. By unanimous consent, the gentleman from Florida could control those 15 minutes.

Mr. GREENWOOD. Mr. Speaker, I ask unanimous consent that the gentleman from Florida (Mr. DEUTSCH) be permitted to control 15 minutes.

The SPEAKER pro tempore. Is there objection to the request of the gentleman from Pennsylvania?

There was no objection.

Mr. DEUTSCH. Mr. Speaker, if I could just inquire, how would we be going in terms of order of speakers?

The SPEAKER pro tempore. The Chair would allow the proponent of the amendment to speak first.